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SATION OF GENE FUNCTION USING DOUBLE STRANDED RNA INHIBITION

(57) Abstract

There is provided a method of identifying DNA responsible for conferring a particular phenotype in a cell which method comprises a) constructing a cDNA or genomic library of the DNA of said cell in a suitable vector in an orientation relative to a promoter(s) capable of initiating transcription of said cDNA or DNA to double stranded (ds) RNA upon binding of an appropriate transcription factor to said or initiating transcription of said cDNA or DNA to double stranged (us) KNA upon binding or an appropriate transcription factor to said promoter(s), b) introducing said library into one or more of said cells comprising said transcription factor, and c) identifying and isolating a particular phenotype of said cell comprising said library and identifying the DNA or cDNA fragment from said library responsible for conferring said phenotype. Using this technique it is also possible to assign function to a known DNA sequence by a) identifying a homologue(s) of said DNA sequence in a cell, b) isolating the relevant DNA homologue(s) or a fragment thereof from said cell, c) cloning said homologue or fragment thereof into an appropriate vector in an orientation relative to a suitable promoter(s) capable of initiating transcription of dsRNA from said DNA homologue or fragment upon binding of an appropriate transcription factor to said promoter(s) and d) introducing said vector into said cell from step a) comprising said transcription factor.

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CHARACTERISATION OF GENE FUNCTION USING DOUBLE STRANDED RNA INHIBITION

The present invention is concerned with characterization or identification of gene function using double stranded RNA inhibition (dsRNAi) and methods of identifying DNA responsible for inducing a specific phenotype in a cell and a method of assigning function to known gene sequences.

It has recently been described in Nature Vol 391, pp.806-811, February 98, that introducing double stranded RNA into a cell results in potent and specific interference with expression of endogenous genes in the cell and which interference is substantially more effective than providing either RNA strand individually as proposed in antisense technology. This specific reduction of the activity of the gene was also found to occur in the nematode worm Caenorhabditis elegans (C. elegans) when the RNA was introduced into the genome or body cavity of the worm.

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The present inventors have utilized this technique and applied it further to devise novel and inventive methods of assigning functions to genes or DNA fragments, which have been sequenced in various projects, such as, for example, the human genome project and which have yet to be accorded a particular function and for use in identifying DNA responsible for conferring a particular phenotype.

30 Therefore, according to a first aspect of the present invention there is provided a method of identifying DNA responsible for conferring a phenotype in a cell which method comprises a) constructing a cDNA or genomic library of the DNA of said cell in an orientation relative to a promoter(s) capable of

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promoting transcription of said cDNA or DNA to double stranded (ds) RNA upon binding of an appropriate transcription factor to said promoter(s), b) introducing said library into one or more of said cells comprising said transcription factor, and c) identifying and isolating a desired phenotype of said cell comprising said library and identifying the DNA or cDNA fragment from said library responsible for conferring said phenotype.

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In a preferred embodiment of the invention the library may be organised into hierarchical pools as described in more detail in the examples provided, prior to step b) such as to include, for example, gene families.

According to a further aspect of the invention there is also provided a method of assigning function to a known DNA sequence which method comprises a) identifying a homologue(s) of said DNA in a cell, b) isolating the relevant DNA homologue(s) or a fragment thereof from said cell, c) cloning said homologue or fragment into an appropriate vector in an orientation relative to a promoter(s) capable of promoting transcription of dsRNA upon binding of an appropriate transcription factor to said promoters, d) introducing said vector into said cell from step a) comprising said transcription factor, and e) identifying the phenotype of said cell compared to wild type.

In each aspect of the invention, the nucleotide or DNA sequence may either be provided in a sense and an antisense orientation relative to a single promoter which has the properties defined above, or alternatively it may be provided between two identical promoters. In both embodiments dsRNA is provided from the transcription initiated from the promoter following binding of its appropriate transcription

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factor.

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The cell according to the invention may be derived from or contained in an organism. Where the cell is contained within an organism, the organism may be adapted to express the appropriate transcription factor. The organism may be any of a plant, animal, fungus or yeast but preferably may be the nematode worm C. elegans, which may be any of a wild type, a nuc-1 or pha-ts mutant of C. elegans or a combination of said mutations. In an alternative embodiment the DNA or cDNA library or the DNA homologue or fragment thereof may, advantageously, be transfected or transformed into a microorganism, such as a bacterial or yeast cell, which may be fed to the organism, which is preferably the nematode worm C. elegans. In this embodiment of the invention the microorganism may be adapted to express the appropriate transcription factor. Preferably, the microorganism is E. coli.

In each aspect of the invention, the DNA library, DNA homologue or DNA fragment may be constructed in a suitable DNA vector which comprises a sequence of nucleotides which encode said transcription factor. Alternatively, said transcription factor is encoded by a further vector. In an even further alternative, the cell or organism may express or be adapted to express said transcription factor. Preferably, any of the vectors used in the method according to the invention comprises a selectable marker which may be, for example, a nucleotide sequence encoding sup-35 or a fragment thereof. The nucleotide sequence may be orientated relative to a promoter such that binding of a transcription factor to the promoter initiates transcription of the DNA into double stranded RNA. Figure 10 illustrates the vectors and the orientation of the DNA sequence which enable double stranded RNA

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production in *C. elegans*. Thus in one embodiment the DNA is located between two promoters on a vector capable of expressing dsRNA upon binding of an appropriate transcription factor to said promoters.

Alternatively, the vector comprises two copies of the DNA sequence organised in a sense and antisense orientation relative to the promoter and which marker is selectable when contained in a pha-1 mutant *C. elegans*. Preferably, the promoters are any of T7, T3 or SP6 promoters and the transcription factor comprises the appropriate polymerase.

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Preferably, the selectable marker comprises a nucleotide sequence capable of inhibiting or preventing expression of a gene in said cell and which 15 gene is responsible for conferring a known phenotype. This nucleotide sequence may be part of or identical to said gene conferring said phenotype, and which nucleotide sequence is itself oriented relative to a suitable promoter(s) capable of initiating 20 transcription of double stranded RNA upon binding of an appropriate transcription factor to said promoter(s). Alternatively, the nucleotide sequence may be a part of or identical to said gene sequence conferring said phenotype, and which nucleotide 25 sequence is such as to permit integration of said suitable or further vector by homologous recombination in the genome of said cell and following said integration said nucleotide sequence is capable of inhibiting expression of said gene sequence conferring 30 said phenotype. In this embodiment said nucleotide sequence comprises stop codons sufficient to prevent translation of said nucleotide sequence following its integration into said genome.

Compounds can, advantageously, in said method be added to said cell or organism for the purposes of

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screening for desired phenotypes, such as for example, resistance or sensitivity to the compound when compared to wild type. The promoters are preferably inducible. The transcription factor may in some embodiments be phage derived, such as for example, a T7 polymerase driven by a phage promoter. However, when C. elegans is utilised a worm specific or tissue specific promoter can be used, such as for example, let858, SERCA, UL6, myo-2 or myo-3. Preferably, the E. coil strain is an RNAaseIII and even more preferably an Rnase negative strain.

A further aspect of the present invention provides a method of generating a transgenic non-human organism comprising an exogenous transcription factor and a transgene comprising a promoter operably linked to DNA fragment which is expressed upon binding of said transcription factor thereto, the method comprising a) providing a first transgenic organism comprising a first construct incorporating DNA encoding an exogenous transcription factor and a second transgenic organism comprising a second construct including at least one promoter operably linked to a desired DNA sequence which is expressed upon binding of the transcription factor of said first transgenic organism thereto b) crossing said first and second transgenic organisms and selecting offspring expressing said desired DNA sequence. In one embodiment said first and second transgenic organisms are generated by transforming said first and second constructs into respective microorganisms for subsequent feeding to the respective organism. Preferably, said second construct comprises said desired DNA sequence in an orientation relative to said promoter so as to be capable of initiating transcription of said DNA to dsRNA upon binding of

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said transcription factor thereto. In this embodiment said second construct comprises two promoters flanking said desired DNA sequence which promoters can initiate transcription of said DNA sequence to dsRNA upon binding of said transcription factor to said promoters. Alternatively, said DNA sequence is provided in a sense and an antisense orientation relative to said promoter so as to produce dsRNA upon binding of the transcription factor to the promoters.. In each of these embodiments the first and/or second constructs may preferably be provided with a reporter gene operably linked to a promoter which is capable of initiating transcription of said reporter upon binding of said transcription factor thereto. Preferably, the reporter gene encodes any of Luciferase, Green Fluorescent protein, β galactosidase or β -lactamase.

The present invention also includes a method of validating clones identified in yeast two hybrid vector experiments which experiments are well known to those skilled in the art and which experiments were first proposed by Chien et al. (1991) to detect protein - protein interactions. The method according to the invention comprises providing a construct including the DNA encoding a protein identified in a two hybrid vector experiment, which construct is such that said DNA is provided in an orientation relative to one or more promoters capable of promoting transcription of said DNA to double stranded RNA upon binding of an appropriate transcription factor to said promoters, transforming a cell, such as a bacterial cell or alternatively transforming an organism comprising said transcription factor with said constructs and identifying a phenotypic change in said cell or organism, which may be C. elegans or the like, compared to wild type. Preferably, the transcription

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factor is inducible in the cell or organism. Once again the DNA sequence may be located between two promoters or in both a sense and antisense orientation relative to a single promoter, as described above. Preferably, the promoter is a phage polymerase promoter and said transcription factor is a RNA polymerase, and preferably T7 polymerases. Also encompassed with the scope of the present invention are vectors used to transform said cells or organisms and the cells or organisms themselves.

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In a further aspect of the present invention there is provided a method of alleviating pest infestation of plants, which method comprises a) identifying a DNA sequence from said pest which is critical either for its survival, growth, proliferation or reproduction, b) cloning said sequence from step a) or a fragment thereof in a suitable vector relative to one or more promoters capable of transcribing said sequence to RNA or dsRNA upon binding of an appropriate transcription factor to said promoters, and c) introducing said vector into the plant.

Thus, advantageously, the method according to the invention provides a particularly selective mechanism for alleviating pest infestation, and in some cases parasitic infestation of plants, such that when the pest feeds on the plant it will digest the expressed dsRNA in the plant thus inhibiting the expression of the DNA within the pest which is critical for its growth, survival, proliferation or reproduction. In a preferred embodiment, the pest may be any of Tylenchulus ssp. Radopholus ssp., Rhadinaphelenchus ssp., Heterodera ssp., Rotylenchulus ssp.,

Pratylenchus ssp., Belonolaimus ssp., Canjanus ssp.,

Meloidogyne ssp., Globodera ssp., Nacobbus ssp.,

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Ditylenchus ssp., Aphelenchoides ssp., Hirschmenniella ssp., Anguina ssp., Hoplolaimus ssp., Heliotylenchus ssp., Criconemellassp., Xiphinemassp., Longidorus ssp., Trichodorus ssp., Paratrichodorus ssp., Aphelenchs ssp. The DNA sequence or fragment thereof according to this aspect of the invention may be cloned between two tissue specific promoters, such as two root specific promoters.

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A further aspect of the invention concerns the vector used in each of the methods of the invention for constructing said library, which vector comprises two identical promoters oriented such that they are capable of initiating transcription of DNA sequence located between said promoters to dsRNA upon binding of an appropriate transcription factor to said promoters. The DNA sequence may, for example, include a multiple cloning site. Preferably, the expression vector comprises a nucleotide sequence encoding a selectable marker. In one embodiment the nucleotide sequence encoding said selectable marker is located between two identical promoters oriented such that they are capable of initiating transcription of DNA located between said promoters to double stranded RNA upon binding of an appropriate transcription factor to said promoters. Preferably, the selectable marker comprises a nucleotide sequence encoding sup-35, for introduction into C. elegans having a pha-1 mutation.

Preferably, the transcription factor comprises either a phage polymerase which binds to its corresponding promoter or a *C. elegans* specific promoter and even more preferably T7 polymerase. Preferably, the vector includes a multiple cloning site between said identical promoters.

In a further aspect of the invention there is provided an expression vector for expressing an

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appropriate transcription factor for use in a method according to the invention which vector comprises a sequence of nucleotides encoding said transcription factor operably linked to suitable expression control sequences. Preferably, the expression control sequences include promoters which are inducible, constitutive, general or tissue specific promoters, or combinations thereof. Preferably, the transcription factor comprises a phage polymerase, and preferably T7, T3 or SP6, RNA polymerase.

A further aspect of the invention provides a selection system for identifying transformation of a cell or organism with a vector according to the invention which system comprises a vector according to the invention wherein said selectable marker comprises a nucleotide sequence capable of inhibiting or preventing expression of a gene in said cell or organism which gene is responsible for conferring a known phenotype. Preferably said nucleotide sequence corresponds to a part of or is identical to said gene conferring said known phenotype, and which nucleotide sequence is itself located between two identical promoters capable of initiating transcription of double stranded RNA upon binding of an appropriate transcription factor thereto. Alternatively, the nucleotide sequence comprises a nucleotide sequence which is a part of or identical to said gene sequence which confers a known phenotype on said cell or organism, and which is such that following integration of said vector by homologous recombination in the chromosome of said cell or organism said sequence inhibits expression of said gene sequence conferring said known phenotype. Preferably, according to this embodiment the nucleotide sequence comprises stop codons sufficient

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to prevent translation of the nucleotide sequence following integration into said chromosome. Preferably, the known gene sequence comprises a sup-35 gene or a fragment thereof which is selectable by identifying offspring growing at a temperature above 25°C following introduction in a pha-1 et123ts mutant C. elegans worm.

In a further aspect of the invention provides said known gene sequence comprises a sup-35 gene or a fragment thereof which is selectable by identifying offspring growing at a temperature above 25°C following introduction of said vector in a pha-1 et123ts mutant *C. elegans* worm. An even further aspect comprises a method of assigning function to a DNA sequence of a multicellular organism which method comprises a) providing i) a construct comprising said DNA fragment cloned between two promoters capable of promoting transcription in said multicellular organism, in a multicellular organism capable of initiating transcription from said promoter; b) identifying the phenotype of said multicellular organism compared to wild type.

The present invention may be more clearly understood by the following examples which are purely exemplary with reference to the accompanying figures, wherein:

Figure 1 is a nucleotide sequence of plasmid PGN1 in accordance with the present invention.

Figure 2 is a nucleotide sequence of plasmid PGN100 in accordance with the present invention.

Figure 3 is a schematic representation of the vectors used and the transformation regime used in the methods according to the present

invention.

Figure 4 is an illustration of an expression vector used in accordance with the invention.

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- Figure 5 is a schematic illustration of the T7 RNA polymerase expression vectors used for transforming *C. elegans*.
- 10 Figure 6 is an illustration of plasmid PGN1.
 - Figure 7 is a diagrammatic representation of an enhanced vector for dsRNA inhibition encoding sup-35 dsRNA.

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- Figure 8 is an illustration of a vector for integration into the genome of *C. elegans*.
- Figure 9 is an illustration of the position of a DNA sequence(s) relative to a suitable promoter to initiate expression of dsRNA from the DNA sequence(s).
 - Figure 10 is a representation of plasmid pGN108.

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- Figure 11 is a representation of plasmid pGN105.
- Figure 12 is a representation of plasmid pGN400.
- Figure 13 is a representation of plasmid pGN401.
 - Figure 14 is a representation of plasmid pGN110.
- Figure 15 is a representation of plasmid pAS2 with forward and reverse T7/T3/SP6 promoters.

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Figure 16 is a representation of plasmid pGAD424 with forward and reverse T7/T3/SP6 promoters.

- Figure 17 is a representation of plasmid pAS2-cyh2-HA+, both T7-final.
 - Figure 18 is a representation of plasmid pGAD424without-FULL-ICE-BOTH-T7.
- Figure 19 (a) is a representation of plasmid pGN205 and(b) is a representation of plasmid pGN207.
- Example A: Construction of an ordered and

 hierarchical pooled cDNA library and applications
 thereof.

A random ordered and pooled library:

promoters, with a multiple cloning site (MCS) in between. The two promoters are orientated towards each other, and towards the MCS. In the presence of T7 RNA polymerase, expressed in E. coli, C. elegans or any other organism, RNA will be produced, starting from the two T7 promoters. As these are oriented in the opposite sense, both strands of RNA will be produced from the DNA inserted (cloned) into the MCS in between the two promoters which results in the generation of double stranded RNA (dsRNA) upon binding of the T7 RNA polymerase thereto.

A C. elegans cDNA library is constructed in the MCS using standard molecular biological techniques. The library is transformed into E. coli, and the resulting E. coli are grown in culture and stored in 96 multi-well plates. At this stage, plasmid DNA can

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be isolated and stored in 96-multi-well plates corresponding to those of the E. coli colonies. Approximately 100,000 colonies are scored. In this way, the library will harbor approximately 5 times the 5 total expressed cDNA variation of C. elegans, which gives the opportunity for low expressed sequences to be present in the library. This will result in approximately 1041 96-well plates. The plates are hierarchical pooled as necessary. For the present 10 pooling of the clones is arranged in a range of 10 to 100. If the hierarchical pooling is per 8 or 12 (numbers are more convenient as 96-well plates have a 8 to 12 grid), this will result in approximately 87 multi-well plates and approximately 8352 wells. If 15 hierarchical pooling is per 96 wells, which is a full plate, this results in approximately 11 plates and approximately 1041 wells. At any stage of the hierarchical pooling, plasmid DNA can be isolated, which would be less elaborate as less plates are used, but will result in a loss of complexity although this 20 should not be the case in the pooling per 12. The pooling of the DNA can also be carried out with the original DNA.

The experiments below describe how the hierarchical pooling should be performed, both for the DNA and for the E. coli library.

An ordered library for RNAi technology, harboring every gene of the C. elegans genome, with applications thereof

As the genome-sequencing project is coming to an end, this information can be used in the application of T7 RNA inhibition technology. Every gene of the C. elegans genome can be cloned using PCR technology. In

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preference, exons will be cloned with a minimal length of 500 bp. If the exons are too small, smaller fragments will be isolated with PCR, or even parts of introns and neighboring exons will be isolated with PCR technology so that at least a sufficient part of 5 the translated region of the gene is cloned. For this, at least 17000 PCR reactions need to be performed. This collection of PCR products will be cloned in a T7 vector as described (two T7 promoters 10 oriented towards each other with a multiple cloning site in between). Every PCR product is cloned independently, or can be used to generate a random library, analogous to the described cDNA library. If every PCR product is cloned individually, the 15 resulting bacteria and plasmid DNA can be pooled in various ways. Firstly, this collection of individually cloned PCR products in the T7 RNAi vector can be pooled randomly, as described in the random library. This pooling can also be done in a more rational way. 20 For instance, the genes of the C. elegans genome can be analyzed using bioinformatic tools (in silico biology). Various genes of the genome will belong to a gene family, or will have homologues in the genome. These members of the gene 25 family will be pooled, or the members, being homologues will be pooled. In this way the total number of about 17000 clones is reduced to a more useable quantity. This library can be used to screen for phenotypes in the methods according to the 30 invention. The resulting phenotype gives a functional description to the gene or gene family or gene homologues of the C. elegans genome. As the library consists of a part of every gene in the genome, this method enables description of the full genome in 35 functional-phenotypic terms. For this the double

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stranded RNA (dsRNA) needs to be introduced in the worm. This introduction of clones alone, or pooled clones, being random pooling or rational pooling can be achieved in several ways as described.

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Example of a vector for the expression of double stranded RNAi

Any vector containing a T7 promoter may be used,
and which contains a multiple cloning site (there are
many commercially available). Primers containing the
T7 promoter and a primer with the reverse
complementary strand, both with the appropriate ends
are designed. These primers can be hybridized, and if
well designed, cloned in the vector of choice. The
minimal sequence for a T7 promoter is
TAATACGACTCACTATAGGGCGA. Although any vector can be
used for the construction of a T7 expression vector
there follows an example of how to achieve this with
the vector pGEM-3zf(-).

- Vector pGEM-3zf(+) (PROMEGA) was digested with HindIII and SalI
- Primers oGN1 and oGN2 were mixed together at a final concentration of 1 $\mu g/30$ μl boiled and cooled slowly to room temperature.
- The primer was ligated into the vector using standard ligation procedures. The resulting vector is pGN1 (shown in Figure 1) and contains two T7 promoters oriented towards each other, and harbors a multiple cloning site in between.

Sequences of oGN1 and oGN2 are:

- oGN1: AGC TGT AAT ACG ACT CAC TAT AGG GCG AGA AGC TT
- oGN2: TCG AAA GCT TCT CGC ATA ATA GTG AGT CGT ATT AC

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Example of the construction of a library

RNA may be isolated from every organism that is sensitive to RNAi. In general the isolated RNA is then copied into double stranded cDNA, and subsequently prepared in suitable vectors for cloning. Several procedures exist and molecular biology kits can be purchased from various firms including promega, clontech, boehringer Mannheim, BRL, etc which enable:

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- isolation of RNA,
- eventually polyA RNA can be isolated (several techniques and kits available)
- first strand synthesis with AMV reverse
- transcriptase, random hexameric primers and/or cligo (dT) primer
 - second strand synthesis with Rnase H, DNA PolymeraseI,
 - flush ends with T4 DNA Polymerase
- 20 addition of an adaptor with T4 DNA ligase.
 - eventually treatment with T4 polynucleotide Kinase
 - cloning of the cDNA into the vector.

The resulting ligation mixture can be considered
as the cDNA library. The ligation contains all cDNA of
the procedure ligated into the vector of interest.
To order the library, the ligation needs to be
transformed into E. coli strains.

30 Application of this E. coli or DNA library

T7 RNA producing strain:

- a standard strain is BL21 (DE3): F-ompT[lon]hsds(r-m-; and E. coli B strain) λ (DE3). Eventually
- variants of BL21 (DE3) can be used, although BL21

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(DE3) pLysS is used.

- any other *E. coli* strain which produces the T7 RNA polymerase, which may be available needs to be constructed. This can be generated easily using a

- phage, which is commercially available, in this case, the $\lambda CE6$ vector (provided by Promega) is used. Almost every $E.\ coli$ strain can be transfected with this phage and will produce T7 RNA polymerase.
 - a RNAseIII mutant E. coli:
- Various strains are in principle available, we chose in a first experiment to use strain AB301-105: rna-19, suc-11, bio-3, gdhA2, his95, rnc-105, relA1, spoT1, metB1. (Kinder et al. 1973 Mol. Gen. Genet 126:53), but other strains may suit better. This strain is
- infected with $\lambda CE6$ and so a T7 producing variant will be constructed.

Wild type *C. elegans* worms can be grown on the bacteria pools. The bacteria is expressing the T7 RNA polymerase. This results in large quantities of dsRNA

- in the gut of the *C. elegans*, which will diffuse in the organism and results in the inhibition of expression. This library can now be used for the screening of several phenotypes. This technique has the advantage that it is a much faster to detect
- relevant genes in certain pathways, than the known C. elegans technology. Moreover, if an interesting phenotype is found, the responsible gene can be cloned easily.
- Using the hierarchical pooling one can easily find in a second screen the relevant clone of the pool. The inserted DNA of this clone can then be sequenced. This experiment results in genetic and biochemical DATA in one step.
- Wild type C. elegans strains can be combined with compounds to screen for phenotype, drug resistance and

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or drug sensibility. The C. elegans strain can be a mutant strain, screening for an enhanced phenotype, reduced phenotype, or a new phenotype. The C. elegans strain can be a mutant strain, and the library screen can be combined with compounds. So one can screen for drug resistance, drug sensibility, enhanced phenotype, reduced phenotype, or a new phenotype. The E. coli strain may be any T7 RNA polymerase expressing strain, like BL21 (DE3), for example, but the formation of double strand RNA may be enhanced by using a special E. coli strain that is RNAseIII negative. RNAseIII recognizes specific loops in dsRNA. Eventually, an E. coli strain can be used that is deleted in RNAses other than RNAseIII or an E. coli can be used that is deleted in one or more RNAses. The expression of the T7 RNA polymerase in most known E. coli strains and constructs which are available to generate T7 RNA polymerase producing E. coli strains, generally comprise an inducible promoter. In this way the production of the T7 RNA polymerase is regulated, and thus the production of the dsRNA. Advantageously, this feature can be used to "pulse" feed the C. elegans worms at specific stages of growth. are grown on the non-induced E. coli strains. When the worm has reached the stage of interest, the T7 RNA production in the bacteria is induced. This allows the studying of the function of any gene at any point in the life cycle of the animal.

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Screening the library for homologues of putative interesting human genes, and assign function to these genes

Hundreds of genes have been isolated in various projects, being genomic projects, differential

expressed arrays, hybridization studies, etc. The described cDNA library can provide a way to validate and or assign function to these genes in a fast and efficient manner. First of all the worm homologue or homologues or the genes need to be identified by bioinformatic tools (in silico biology). PCR primers are developed and the cDNA fragment is isolated using PCR technology. PCR can be performed on the hierarchical pools. The positive pool or individual wells harboring the bacteria that has the appropriate cDNA is fed to *C. elegans* and the phenotype is scored.

PCR can be performed on cDNA isolated from *C*. elegans. The resulting DNA can be cloned in the T7 vector and transformed in the dsRNA producing *E*. coli on which the *C*. elegans worms are then fed. Depending on which way is faster and more reliable a choice needs to be made.

If the gene belongs to a gene family, the worm may need to be fed on a mixture of bacteria. Each of them harboring a part of the member of the gene family. E. coli strains, growth conditions, combinations with compounds can be performed as described above.

If the library rational is used, in which all the genes of *C. elegans* are cloned in a organized and structured way, the *C. elegans* homologue and eventually the other homologues, orthologues, and members of the gene family can be traced back easily in the library using a silico biology. No PCR is involved in this step, and the bacteria and or DNA can be isolated on which the worm will be grown.

Examples

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The idea of the series of experiments was to test

both the RNAi vector and the various E. coli strains that were constructed

- Construction of a test plasmid 5 Any cDNA that gives a clear phenotype in the worm when knocked-out, or used in a RNAi experiment can be used. It is known that unc-22 is a good candidate, but may other genes are possible. We opted for a sensitive system that can be used at a later stage. 10 The system was tested with sup-35 in a pha-1 background. Exon 5 of the sup-35 was isolated by PCR and cloned in the T7 promoter vector pGN1. The resulting vector was designated pGN2. pha-1 (e2123) mutant worms cannot produce offspring at temperatures higher than 25°C. This is due to a developmental 15 problem in embryogenesis. When sup-35 is knocked-out, or inhibited in this strain, offspring may grow at this temperature. Combination of pha-1 mutant worms and sup-35 RNAi is a good system to validate the 20 various options.
- Testing the RNAi using an E. coli strain that produces dsRNA. - pGN2 was introduced in E. coli strain BL21(DE3) and 25 T7 RNA polymerase was induced with IPTG. C. elegans worms (pha-1 (e2123)) were inoculated on this bacteria, and grown at the restricted temperature of 25°C. As this mutant is an embryonic mutant at this temperature, no offspring will be observed. If the 30 sup-35 gene is efficiently inhibited by the dsRNA present in the E. coli, offspring will be observed. - pGN2 was introduced in E. coli strain AB301-105(DE3) and T7 RNA polymerase was induced with IPTG. C. elegans worms (pha-1 (e2123)) were inoculated on this 35 bacteria, and grown at the restricted temperature of

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25°C. As this mutant is an embryonic mutant at this temperature, no offspring will be observed. If the sup-35 gene is efficiently inhibited by the dsRNA present in the *E. coli*, offspring will be observed.

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3) Improving the worm strain for better uptake of dsRNA.

Before plating the pha-1 *C. elegans* on the *E. coli* strain that produce the double stranded sup-35 RNA. The worm was mutagenised with EMS (Methane sulfonic Acid Ethyl). The offspring of this mutagenised worm is then plated on the bacteria. The worm that feed on this bacteria give larger offspring which has a mutation that results in an improvement of dsRNA uptake, and can be used for further experiments.

Stable integration of the dsRNA producing vector into the genome of the T7 RNA polymerase producing worm

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An E. coli vector can be constructed harboring the following features; Two T7 promoters directed towards each other, with a restriction site or a multiple cloning site in between. Furthermore, the vector may contain the C. elegans sup35 genomic DNA, engineered in such a way that it contains several stopcodons at various intervals, so that no full length protein can be expressed form the sup35 genomic DNA fragment as illustrated in Figure 8. Any cDNA or cDNA fragment can be cloned in the multiple cloning site between the two T7 promoters. When this vector is introduced in a C. elegans strain which expresses T7 RNA polymerase, the cDNA or DNA fragment cloned between the two T7 promoters will be transcribed, generating dsRNA from the cloned fragment.

The vector is designed to be used in pha-1 (e2123) mutant worms expressing T7 RNA polymerase. The expression of the T7 RNA polymerase may be constitutive or regulated, general or tissue specific. These pha-1 (e2123) worms cannot produce offspring at temperatures higher than 25°C, which is due to a development problem in embryogenesis. When sup-35 is inhibited or knocked-out in this stain, offspring may grow at this temperature.

10 When the vector is introduced in the worm, the vector may integrate by homologous recombination (Campbell-like integration). It has been shown that homologous recombination occurs in C. elegans, although at low frequencies (Plasterk and Groenen, EMBO J. 11:287-290, 1992). Homologous recombination 15 at the sup35 gene will result in a knock-out of the gene as the two resulting sup-35 genes will harbor the stopcodons. The resulting worm, and its offspring, if this recombination happens in the eggs, will have a 20 copy of the vector integrated in the genome. This can be selected as only the worms for which the sup-35 has been knocked-out will have offspring at temperatures higher than 25°C. Furthermore, the resulting worm will stably produce double stranded RNA from the DNA 25 fragment cloned between the two T7 promoters. This worm can now be considered as a stable transgenic worm strain with a reduction of function of the gene, from which a fragment has been cloned between the two T7 promoters.

The DNA may be delivered to the worm by several techniques, including injection, ballistic transformation, soaking in the DNA solution, feeding with bacteria. New and other methods that increase the transformation efficiencies can be considered.

The target C. elegans strain may in addition,

have other mutations than the pha-1 (e2123) mutation, and may express other genes than T7 RNA polymerase.

EXAMPLE B: a Yeast two-hybrid-RNAi vector

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A yeast two hybrid vector can be constructed harboring the two T7 promoters. The vectors can be designed to replicate both in yeast and in E. coil. In general cDNA libraries for the yeast two hybrid 10 system are made in the Gal4 or LexA vectors. The library is constructed in vectors having the activation domain of one of these genes. A vector can be constructed that can still perform in the yeast two hybrid screen but which also contains two T7 15 promoters orientated towards each other, with a cloning site therein between. The order of the sequences in the plasmid will then be "plasmid backbone, (GAL4-T7), MCS, T7, backbone". A C. elegans cDNA library constructed in this vector can be used as a standard yeast two hybrid library in an experiment to isolate interacting proteins with a given protein. Once a clone is isolated, the plasmid can be introduced in an E. coil strain expressing the T7 RNA polymerase, and hence will produce dsRNA of the 25 cloned fragment. The bacteria producing this dsRNA can be fed to the worm and phenotypes can be scored. As in the previous example, this validation procedure for a newly isolated yeast two hybrid clone is remarkably shorter than the standard procedure, which requires PCR and/or cloning steps, RNA experiments and/or knock-out experiments. In most cases isolated clones are sequenced first, and on the basis of the sequence, a decision is made to continue with further experiments. In the present invention every isolated clone can easily be introduced into the appropriate E.

coli and fed to the worm. Validation is then performed by phenotype analysis.

To apply this procedure a yeast two hybrid was performed using a known gene as bait and the newly 5 constructed library as the target. Proteins coded by the clones in the target that interact with the bait protein, will result in positive yeast clones expressing the reporter molecule such as can be observed by LacZ staining with X-gal. The plasmid 10 coding for the target protein is isolated directly from the yeast strain and introduced in E. coli. The E. coli is T7 RNA polymerase producing E. coli. In this case, double stranded RNA is produced from the DNA cloned in the multiple cloning site of the vector. 15 When this dsRNA is fed to the worm using the methods described previously, the gene has inhibited in the worm, resulting in a particular phenotype. - This yeast two hybrid vector can advantageously be used to construct an ordered and hierarchically pooled 20 library as described in the previous example. - A yeast strain can also be constructed that conditionally produces T7 RNA polymerase. After yeast two hybrid experiments, the expression of the T7 polymerase could be induced, resulting in the 25 production of dsRNA in the yeast cell. Consequently

Construction of a T7 RNA polymerase producing strain, and applications thereof

on yeast.

the yeast could be fed to the worm. Evidence is

available showing that the C. elegans worms can feed

A C. elegans strain can be constructed that expresses T7 RNA polymerase. The expression can be general and constitutive, but could also be regulated

under a tissue specific promoter, an inducible promoter, or a temporal promoter or a promoter that harbors one of these characteristics or combination of characteristics. DNA can be introduced in this C.

- elegans strain. This is done either by injection, by shooting with particles, by electroporation or as aforementioned by feeding. If the DNA is a plasmid as described in the previous examples, i.e. a plasmid harboring a cloned cDNA fragment or a PCR fragment
- between two flanking T7 promoters, then dsRNA of this cDNA or PCR fragment is formed in the cell or whole organism resulting in down regulation of the corresponding gene. The introduced DNA can have an efficient transient down regulation. The introduced
- DNA can form an extrachromosomal array, which array might result in a more catalytic knock-out or reduction of function phenotype. The plasmid might also integrate into the genome of the organism, resulting in the same catalytic knock out or reduction
- of function phenotype, but which is stably transmittable.
 - Plasmid DNA harboring a cDNA or a part of a cDNA or an EST or an PCR fragment of *C. elegans* cloned between two T7 promoters as described in Examples A) and B)
- can be introduced in the T7 RNA polymerase worm, by standard techniques. Phenotypes can be analysed -DNA from an ordered and pooled library as in Example A) can be introduced in the T7 RNA polymerase worm, by standard techniques (injection, shooting). Phenotypes
- can be analysed. With the hierarchical pool, the original clone can be found easily.
 - The same procedure can be performed with a mutant worm expressing the T7 RNA polymerase. Screening for enhanced, reduced or new phenotypes.
- 35 The procedure can be used to enable screening of

compounds. Screening with either a wild-type strain or a mutant strain for enhanced or new phenotypes.

- The DNA could be introduced in the worm by new methods. One of which is the delivery of DNA by E.
- 5 coli. In this case the hierarchical pooled library is fed to the animal. To prevent digestion of the E. coli DNA in the gut of the nematode, preferentially a DNAse deficient C. elegans will be used, such as nuc-1 (e1392). This procedure would be one of the most
- interesting as it would be independent of transformation efficiencies of other techniques, and generally faster and less labourious.
 - 2) Putative enhancements of the method.
- A vector is designed, so that it harbors the sup-35 cDNA or a part of this cDNA, cloned in between two T7 promoters. The rest of the vector is as described in Examples A) and B). This vector can be introduced into a pha-1ts mutant *C. elegans*. A temperature selection
- system exists in this case and only those worms which have taken up the DNA and express the double stranded sup-35 RNA will survive at restricted temperatures. The hierarchical pooled library can be delivered by any method described above.
- 25 The vector can be used to construct a library that is introduced in a T7 RNA polymerase expressing E. coli. In this case we have an analogous screening as in part A) with an additional screening for worms where the dsRNA of sup-35 is active.
- The DNA and or dsRNA of sup-35 could be delivered on a different plasmid. For the feeding, both DNA feeding (Example C) or dsRNA feeding Example A) and B), this means that the two plasmids could be present in one bacterium, or that the worm is fed on a mixture of

bacteria, one of which harbors the sup-35 construct.

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Example of the construction of a T7 RNA producing C. elegans

To produce T7 RNA polymerase in the worm, several possibilities are possible. The T7 polymerase can be expressed under various promoters, being inducible promoters, constitutive promoters, general promoters and tissue (cell) specific promoters, or combinations of those. Examples of these promoters are the

heatshock promoter hsp-16, the gut promoter ges 1, the promoter from cet858, but also the promoter of dpy 7 and the promoter element GATA1. In this example the T7 RNA polymerase is expressed under the control of the hsp-16 promoter that is available in the pPD49.78

vector. The T7 RNA polymerase is isolated as a PCR product using the primers of GN3 an GN4.

The resulting PCR product is digested with NheI and NcoI, as is the vector in which we want to clone, being the Fire vector pPD49.78. The resulting vector is pGN100 illustrated in Figure 2 oGN3: CAT GGC AGG ATG AAC ACG ATT AAC ATC GC oGN4: ATG GCC CCA TGG TTA CGG GAA CGC GAA GTC CG pGN100 is included.

The vector is introduced into the worm using standard techniques, such as micro injection, for example.

The following strains were then constructed:

-Wild-type (pGN100)

-nuc-1 (e1392) (pGN100)

-pha-1 (e2123) (pGN100)

-pha-1; nuc-1 (pGN100)

All of these strains are able to produce T7 RNA polymerase when temperature induced or alternatively

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by metals such as application of heavy cadmium or mercury. The procedure for temperature induction is to shift the animal to a temperature of 30-33°C for at least one hour, then the animal can be shifted back to standard

temperatures (15-25°C).

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The wild type strain producing T7 RNA polymerase can be used for the production of any RNA in the worm. More specifically, the plasmids from the described libraries can be introduced in these worms, and phenotypes can be scored.

The nuc-1 mutant worm will be used to introduce DNA via bacteria on which the worm feed. As the nuc-1 worm does not digest the DNA, the plasmid DNA can cross the gut wall. If taken up by the cells that produce the T7 RNA polymerase, dsRNA will be produced thus inhibiting the gene from which the RNA was transcribed.

The pha-1 mutant strain that produced T7 RNA polymerase can be used to enhance the procedures as described above. DNA can be introduced by shooting, micro injection or feeding. More specifically this strain can be used for the vectors that produce dsRNA from sup-35 and from the gene of interest, the latter can be a PCR product, a cDNA, or a library as described.

The pha-1; nuc-1 mutant producing T7 RNA polymerase can be used for the bacterial delivery of the DNA. DNA will preferentially be the plasmid that produce dsRNA from both sup-35 and the gene of interest. The worm strain will preferentially produce the T7 RNA polymerase in the gut. Delivery will preferentially happen by feeding the worm on bacteria harboring the plasmid.

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Application of the RNAi technology in plants

Nematodes are responsible a large part of the damage inflicted on plants and more particularly to plants used in the agricultural industry. The RNAi procedures according to the invention can be applied to plants to prevent these parasitic nematodes from feeding longer. In a first step, a DNA fragment is isolated from the parasitic plant nematode that is critical for the animals survival or growth, or to feed or to proliferate. Any gene from which the expression is essential is suitable for this purpose.

A part of this gene, an exon or cDNA is cloned. This DNA fragment can be cloned under the influence of a tissue specific promoter preferably a root specific promoter even more preferably between two root specific promoters. The DNA of the cloned gene under the control of the root specific promoter can be introduced in the plant of interest, using plant transgenic technology. For every parasitic nematode, a different piece of DNA may be required and likewise for every plant race, a different promoter will be needed.

The root will produce RNA or dsRNA from the introduced piece of DNA when root specific promoter is utilised. As the nematode feeds on the plant, the RNA and/or dsRNA will be consumed or ingested by the nematode. The RNA and/or dsRNA can enter the cells of the nematode and perform its inhibitory action on the target DNA. Depending on the nature of the cloned DNA piece of worm, the nematode will not be able to survive, to eat, proliferate, etc in any case preventing the animal of feeding longer on the plant, and thus protecting the plant.

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Construction of a T7 RNA polymerase producing C. elegans

To produce a T7 RNA polymerase or other RNA

polymerases in animals, and more particularly in
nematodes and most particularly in C. elegans, several
possibilities can be envisaged. The T7 RNA polymerase
can be expressed under various promoters. These
promoters may be inducible promoters, constitutive

promoters, general promoters, tissue specific
promoters, or combinations of those.

Example1:

Construction of a basic vector for expression of T7 polymerase in C. elegans

The T7 polymerase coding sequence was PCR amplified from \(\lambda \) CE6 (Novagen, Madison, USA) using the primers oGN26(ATGGAATTCTTACGCGAACGCGAAGTCCG) and 20 oGN46(CTCACCGGTAATGAACACGATTAACATCGC), using standard procedures (PCR, A practical A practical approach, 1993, Ed. J. McPherson, et al, IRL Press). The resulting DNA fragment encoding for the T7 RNA polymerase was digested with AgeI and EcoRI and 25 inserted into the Fire vector pPD97.82 digested with AgeI and EcoRI. The resulting construct encodes for an open reading frame of T7 RNA polymerase in fusion with the SV40 large T antigen nuclear localization signal (NLS) with amino acid sequence MTAPKKKRKVPV. This 30 nuclear localization signal sequence is required to translocate the T7 RNA polymerase from the cytoplasm to the nucleus, where it is able to bind to its specific promoters, designated T7 promoters. Upstream of the coding sequence for the T7polymerasefusion 35 protein is a minimal promoter (myo-2) preceded by a

multiple cloning site (MCS) in which several C.
elegans promoters can be inserted. This plasmid
(pGN105 shown in Figure 11) is a basic T7 RNA
polymerase plasmid which enables the expression of
T7polymerase in C. elegans. Derivatives of this
plasmid wherein promoters are cloned into the multiple
cloning site, allow for the inducible, constitutive,
general and tissue specific expression of T7 RNA
polymerase in C. elegans, as expression will be
regulated by the promoter cloned in the multiple
cloning site.

Although not restricted to these examples, for the following promoters it is known that they induce expression in the following tissues.

15 let-858 (ubiquitous expression), myo-2 (pharynx
expression), myo-3 (body wall muscles), egl-15 (vulval
muscles), unc-119 (pan-neuron).

Example 2:

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20 <u>Construction of a vector for expression of T7 RNA</u>
polymerase in C. elegans muscle tissue.

The T7 RNA polymerase coding sequence was PCR amplified from_\(\)\(\text{CE6}\) using the primers oGN43 (GCCACCGGTGCGAGCTCATGAACACGATTAACATCGC) and oGN44 (CACTAGTGGGCCCTTACGCGAACGCGAAGTCCG) digested with AgeI/SpeI and inserted in the pGK13 vector digested with AgeI/SpeI. (This vector contains the strong SERCA promoter which drives expression in the pharynx, the vulval muscle, the tail and the body wall muscle). A nuclear localization signal (NLS) of SV40 large T antigen was inserted in front of the T7 polymerase coding sequence by insertion of two overlapping oligo's oGN45 (CCGGATGACTGCTCCAAAGAAGAAGCGTAAGCT) and oGN46 (CTCACCGGTAATGAACACGATTAACATCGC) into the

SacI/AgeI restriction sites. The resulting construct was called pGN108 as shown in Figure 10. Introduction of this plasmid into C. elegans results in the expression of T7 RNA polymerase in the pharynx, vulva muscle, tail and body wall muscles.

To test expression and functionality of T7 RNA polymerase in C. elegans under the regulation of the SERCA promoter, pGN108, which encodes the T7RNA polymerase under the control of the SERCA promoter was 10 injected into C. elegans. A test vector was coinjected. This test vector encodes for GFP under the control of a T7 promoter (pGN401 in Figure 13). The plasmid pGN401 was constructed by inserting two overlapping oligo's oGN41 (CCCGGGATTAATACGACTCACTATA) 15 and oGN42 (CCGGTATAGTGAGTCGTATTAATCCCGGGAGCT) in the SacI/AgeI opened Fire vector pPD97.82. generating a T7 promoter. Furthermore a selection marker was coinjected to select for transformants (rol6, pRF4). The latter selection vector pRF4 is well known to person 20 skilled in the art. Transgenic F1 could easy be isolated as they display the rol 6 phenotype. These transgenic C. elegans all expressed GFP in the pharynx, the vulval muscle, the tail and the body wall muscle. This data show clearly that the T7 RNA 25 polymerase is functionally expressed under the regulation of the SERCA promoter, and that the expressed T7 RNA polymerase binds to the T7 promoter present in pGN401 and initiates transcription of the GFP gene, which is then functionally expressed, 30 resulting in fluorescence in the muscle tissues where SERCA is inducing the expression of the T7 RNA

polymerase.

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expression of T7 polymerase in C. elegans

Example 3: Construction of a vector for ubiquitous

5 The NLS-T7 RNA polymerase fusion gene was isolated from pGN108 with XmaI/Bsp1201 and cloned into the Fire vector pPD103.05 digested with XmaI/Bsp120I.This results in a vector wherein the T7 RNA polymerase is cloned under the regulation of the let858 promoter. This specific promoter enables the expression of T7 RNA polymerase in all tissues. The resulting plasmid was named pGN110 (Figure 14).

Example 4:

Construction of a vector for T7 RNA polymerase mediated expression of DNA fragments, genes, and CDNA's under the control of a T7 promoter.

The Fire vector pPD97.82 was digested with SacI / 20 AgeI and a T7 promoter sequence was generated by insertion of two overlapping oligo's oGN41 (CCCGGGATTAATACGACTCACTATA) and oGN42 (CCGGTATAGTGAGTCGTATTAATCCCGGGAGCT) into the SacI/Age/ restriction endonuclease sites. This construct (pGN400 25 Figure 12) contains a GFP open reading frame cloned between SacI and EcoRI restriction endonuclease sites under the regulation of the T7 promoter. Any gene, cDNA, or DNA fragment can be cloned in this vector by deleting the GFP gene as a AgeI/ SacI fragment and 30 cloning the DNA fragment of interest into the vector. Preferentially the DNA fragment of interest can be obtained by PCR amplification, inserting the SacI/AfeI sites in the primers. The resulting DNA fragment after PCR amplification is the digested and the GFP gene in 35 pGN400 is replaced by the amplified DNA fragment.

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Every vector that contains a T7 promoter could be used for the purpose of T7 RNA polymerase induced expression in C. elegans, such as the commercially available pGEM vectors and the pBluescript vectors. This is clearly shown by the pGN401 vector which expresses GFP under the regulation of the T7 promoter in a transgenic C. elegans which expresses T7 RNA polymerase.

The use of pGN400 has the advantage that the vector includes a 3'UTR fragment from unc-54 which enhances the transcription or stability of the RNA.

Generation of permanent, tissue specific "pseudo knock-out" RNAi C. elegans lines

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At present, gene knock outs in C. elegans are obtained after random, large scale mutagenesis and PCR base sib-selection. This method is bulky, very time consuming and tedious. It has been described that introducing double stranded RNA into a cell results in potent and specific interference of expression of endogenous genes. In C. elegans gene expression can be down regulated by injection of RNA into the body cavity of the worm, soaking the worm in a solution containing dsRNA or feeding E. coli that express dsRNA corresponding to the gene of interest. C. elegans cells have the ability to take in dsRNA from their extracellular environment. It has been reported that mRNA is the target of this ds RNA mediated genetic interference (Montgomery and Fire 1998). It is also suggested that the targeted RNA is degraded in the nucleus before translation can occur. Although the RNAi mediated reduction of gene expression can be passed on to the next generations, heritability is poor and the effect is rapidly lost during further

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offspring. This is probably due to a continued decrease of the dsRNA pool. We propose here a method to construct C. elegans lines with a permanent, inheritable, RNAi phenotype. The method encompasses the generation of transgenic C. elegans lines by introducing plasmids containing cDNA fragments of the target gene in the sense and antisense orientation under control of a worm promoter or by transcription of an inverted repeat of the cDNA from a single construct. Alternatively, ds RNA can be transcribed from a vector harboring a cDNA flanked by two T7 promoters in a C. elegans strain that expresses T7 polymerase. The result is a transgenic worm with an heritable stable "pseudo knock-out" phenotype. The expression of the cDNA or the T7 polymerase can be general and constitutive but could also be regulated under a tissue specific promoter. In contrast to RNAi induced by external ds RNAi (injected, soaked or feeded) this method would enable to obtain conditional, tissue specific inhibition of gene expression.

Inhibition of unc-22 expression by RNA interference results in a "twitching" phenotype.

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Unc 22 cDNA (exon 22) was cloned in sense and antisense orientation in pPD103.05. (A. Fire nr L2865) containing the let 858 promoter that is capable of expressing RNA sequences in all tissues. The resulting plasmids were named pGN205 (Figure 19a) and pGN207 (Figure 19 b). These constructs were introduced into C. elegans together with a selectable marker (rol-6; GFP). Transgenic F1 individuals (expressing rol-6 or GFP) showed a "twitching" phenotype indicating that RNAi could be mediated by endogenous transcription of

RNA from transgenic DNA. The RNAi phenotype cosegregated with the selectable marker into further offspring. This resulted in the generation of C. elegans lines with permanent RNAi phenotype.

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Generation of stable lines T7 RNA polymerase lines and generation of dual transgenic worms.

An expression system in C. elegans based on an exogenous RNA polymerase demands two plasmids. One is encoded for the RNA polymerase under the control of a specific promoter, while the other plasmid encodes for the DNA fragment to be expressed, under the regulation of the T7 promoter. In the case of semi stable RNAi also designated pseudo stable knockouts, the DNA of interest is cloned between two T7 promoters so that dsRNA can be produced.

As the T7 RNA polymerase expression system is known to be a high expression system this will result in problems to generate dually transgenic animals. If the gene to be expressed in the C. elegans nematode is toxic, this will result in lethal effects and hence in the construction of a C. elegans without highly regulated stable expression of the gene of interest. If the gene of interest is essential for the survival of the organism, RNAi with a DNA fragment from this gene will also result in lethal effects, so that pseudo- stable knockouts are not possible.

To overcome this problem the present inventors have designed a system consisting of two transgenic animals. The first animal is transgenic for the T7 RNA polymerase, This T7 RNA polymerase can be expressed in all cells or specific cells or tissues as has been shown in previous examples. The second transgenic animal is transgenic for the DNA fragment of interest.

This can be a gene or cDNA linked to a T7 promoter, or if one wants to perform RNAi a DNA fragment of such gene cloned between two T7 promoters.

Both transgenic animals are viable and do not show any aberrant phenotypes. This is because the T7RNA polymerase expressed in the first transgenic organism is not toxic for the organism, even if expressed at relative high levels. In the second transgenic organism, the gene of interest is not expressed or the dsRNA is not produced as these transgenic animals do not contain the T7 RNA polymerase.

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Expression of the gene or cDNA of interest or RNAi with a DNA fragment can now be obtained by mating the two transgenic animals. The offspring of these are dually transgenic and express the gene of interest or express dsRNa of the DNA fragment of interest.

To generate sufficient males in such a mating, one of the transgenic animals males can be a C. elegans mutant with a phenotype favouring generation of males. An Example of such a mutant is him-5. Preferentially such a mutant will be used to make a C. elegans transgenic for T7 RNA polymerase, while the hermaphrodite harbors the DNA fragment under the regulation of the T7 promoter.

To select efficiently for the dual transgenic offspring a second transgene can be introduced in the second transgenic animal. This transgene contains a reporter gene under the regulation of the T7 promoter. The reporter gene can be GFP, luciferase, Beta galmactosidase, or beta-lactamase. an example of such a ttransgene are the vectors pGN400 and pGN401.

To obtain inducible, tissue specific expression of a transgene in C. elegans we can make male stock (i.e. him-5) carrying the T7 polymerase construct

under the control of different C. elegans promoters that enable tissue specific expression such as). This males can be crossed with hermaphrodites carrying the gene of interest under the control of a T7 promoter.

Furthermore, the transgenes can be integrated into the genome of the animal. Methods to generate stable integration of a plasmid into the genome of the animal have been described (Methods in cell biology, Vol. 48, 1995, ed. by epstein and Shakes, academic press) and involve radiation of the animal. This can be done for both animals animals, but preferentially, the animals expressing the T7 RNA polymerase are subject to such traetment. This result in a collection of C. elegans nematodes that stably express T7 RNA polymerase under the control of various promoters. examples of such promoters are the myo-2 (pharynx expression), myo-3 (body wall muscles), egl-15 (vulval muscles), unc-119 (pan-neuron), SERCA (muscles), let858 (all cells) ges-1 (gut).

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Construction of RNAi T7 promoter yeast two hybrid vectors

In most two-hybrid experiments a cDNA library is cloned in plasmid pGAD424 (Figure 16) which has been engineered with additional restriction sites in the polylinker such as a Ncol site (Clontech). This library allows for screening of binding proteins in a yeast two hybrid experiment. We constructed a new yeast two hybrid vector with the same possiblilities to perform yeast two hybrid, but which contain two addditional T7 promoters, so that the vector can be used for T7 RNA polymerase induced pseudo-stable knock-outs. For this we inserted a forward T7 by using

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a T7- linker (consisting of the following primers aattettaatacgactcactatagggec and catgggccctatagtgagtcgtattaag) into the EcoRI-Ncol site of pGAD424. The resulting vector was designated pGAD424-without-FULL-ICE-both-T7. Care was taken to eliminate stop codons and using maximal polylinker compatible amino acids. We adopted the same strategy for the reverse T7 (consisting of both primers gatccgtcgacagatctccctatagtgagtcgtattactgca and gtaatacgactcactatagggagatctgtcgacg) with BamH1 and Pst1. To avoid loss of SalI, we included this site in the primer.

The SalI site is important as most libraries are cloned in this site, adapters are available. This makes the newly constructed vector compatible with existing vectors.

pAS2 with with forward and reverse T7/T3 and or Sp6

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An analogous yeast two hybrid vector was constructed based on pAS2 (Clontech). By partial EcoRV digestion we were able to remove a significant part of the cyh2 gene. The right construct can be isolated and checked by a restriction digest with BglII. this restriction site is present in the EcoRV fragment of PAS2 to be eliminated. This elimates the cyh2 gene which is slightly toxic gene and involved in growth retardation. This gene is non-essential for the performing of RNAi and Yeast two hybrid experiments. After the elimination of the EcoRV fragment, The EcoRI restriction site which is located between the DNA sequence encoding for GAL4DB and HA (epitope) becomes unique for the plasmid, and can be used to subsitute HA with a T7 promoter containing linker.

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This ensures persistence of all restriction sites, allowing both in frame cloning and compatibility with previous vectors and pGAD424. We used the following linker (primers: aattcttaatacgactcactatagggca and

tatgccctatagtgagtcgtattaag) using EcorI and Ndel cloning sites. We adopted the same strategy for the reverse T7 (primers:

gatecgtegacagatetecetatagtgagtegtattactgca catgggeectatagtgagtegtattaag and

gtaatacgactcactatagggagatctgtcgacg) with BamH1 and Pst1. To avoid loss of Sall we included it in the primer. The resulting vector was designated pAS2-cyh2-HA+both T7-final.

Having the T7 promoter (or alternatively the T3, or SP6 promoter) in pGAD424 allows to go quickly from interacting protein to RNAi and assigning function to the isolated DNA fragment. An additional advantage is the ability to make by in vitro transcription coupled to in vitro translation (There is an ATG in frame with either GAL4DB or GAL4AD) labeled protein which can be used for in vitro controls (e.g. pull down assays) of the actual protein-protein interaction.

The sequences of the plasmids produced and the SP6 and T3 polymerase are identified in the Sequence Listing provided below:

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SP6 DNA-dependent RNA polymerase:

SEQUENCE ID NO. 1

swissprot accesion number P06221

protein sequence:

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1 mqdlhaiqlq leeemfnggi rrfeadqqq iaagsesdta wnrrllseli apmaegiqay
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T3 DNA dependent RNA polymerase:

SEQUENCE ID NO. 2

swissprot accession number P07659

protein sequence:

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pGN108:

SEQUENCE ID NO. 3

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tree the titigactic telestate case a casectcl caata tatta ctagggag acgaggagg accttate gaa cgtcg catgag ccattett ctt cttattat catctct ctcacacatta at a constraint constraintgageegaaaaaacateegtagtaagtetteetttaageegacaettttagacageattegeegetagttttgaagtttaaattttaaaaaataaaaattag to the against a capacity of the control of the cagittgttttaaaggaaaaacacgaaaaaagaacactatttatctttcctccccgcgtaaaattagttgttgtgataatagtgatccgctgtctatttgcact eggetetteacacegtgetteeteteacttgacecaacaggaaaaaaaaacateacgtetgagaeggtgaattgeettateaagagegtegtetetttea ctccagcagcaccacaatcatcttcgtgtgttctcgacgattccttgtatgccgcggtcgtgaatgcaccaccattcgacgcgcaactacaccaccaccactcut and the second content of the second content oactive get get attacta cacegicate get the transfer of the tr $caccactect gac g \pi t ctacc the tight the city of the case of the$ tictagigitta atticegea taa attiggacetaa aattigggittitigicate attitica acaa gaa ategigaa aateet gitigittiegea attitietitie aa aaa ataa aattigi gaa aacacgaaatatatggtaatttcccgaaatattgagggtctcgccacgatttcagtcacagtggccaggatttatcacgaaaaaagttcgcctagtctcacat trccggaaaaccgaatctaaattagtttttgtacattttgaacaaaaaatcgagacatccctatagtttcgcaattttcgtcgcttttctccaaaaatga cagtetagaattaaaattegetggaaetgggaeeatgatatetttteteeeegttttteattttatttttatataeaetggattgaetaaaggteaeeaeeaeeg ccagigigigecatateaeaeaeaeaeaeaeaeaeatgiegagaittiatgigitateeetgettgattiegtteegtigietetetetetetatteatetttig agccgagaagetccagagaatggagcacacaggatcccggcgcgcgatgtcgtcgggagatggcgccgcctgggaagccgccgagagatatca gggaagategtetgattteteeteggatgeeaeeteatetetegagttteteegeetgttaeteeetgeegaaeetgatattteee

pGN105:

SEQUENCE ID NO. 4

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pGN400:

SEQUENCE ID NO. 5

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actegagtaatatttaaatttteageatetegegeeegtgeetetgaettetaagteeaattaetetteaaeateeetaeatgetettteteeetgtgeteeeaeee ataaattttttttgaaacatcatagaaaaaccgcacacaaaataccttatcatatgttacgtttcagtttatgaccgcaatttttatttcttcgcacgtctgggc ggggtttcccctattgtttgtcaagagttcgaggacggcgttttcttgctaaaatcacaagtattgatgagcacgatgcaagaagatcggaagaagg at accasa a cata act g tttcct act agt cgg ccg tacggg cccttt cgt ctcg cgcgttt cgg tgatgacggt gaaaacct ct gacacat g cagct cccg gacacat g cagct g cgagacggicacagctigicigiaagcggaigccgggagcagacaagcccgicagggcgcgicagcgggigitggcggggigicggggciggctia actat tgcgg catcagag cagattg tactgag ag tgcaccatat tgcgg tgtgaaataccgcacagattgcgtaag gagaaaataccgcatcagg cggcaccatattgcgg tgtgaaataccgcacagattgcgtaag gagaaaataccgcacagattgcgtaag gagaaaataccgcacagattgcgtaag gagaaaataccgcacagattgcgcaccatattgcgg tgtgaaataccgcacagattgcgtaag gagaaaataccgcactagattgcgcaccatattgcgg tgtgaaataccgcacagattgcgtaag gagaaaataccgcacagattgcgcaccatattgcgg tgtgaaataccgcacagattgcgtaag gagaaaataccgcacagattgcgcaccatattgcgg 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gag ta cae ta the tattete agaat gac tt gg tt gag ta cae ta the tattete agaat gac tt gg tt gag ta cae ta the tattete agaat gac tt gg tt gag ta cae ta the tattete agaat gac ta ta cae ta the tattete agaat gac ta ta cae tacagicacagaaaagcatcttacggaiggcaigacagtaagagaattaigcagigcigccataaccaigagigataacacigcggccaacttacticiga gigagegigggiciegeggiateatigeageaetggggeeagatggtaageeeteeegtategtagtiatetaeaegaeggggagteaggeaaetat actic attitta a titta a a aggai citaggi ga agai cottitti gata a tot cat ga coa a a at cootta a citaggi ga ga titto ga coa cataga cootta ga cootta a citaggi ga ga titto ga coa cataga cootta ga cootta a citaggi ga ga titto ga coa cataga cootta ga cootta a citaggi ga ga titto ga coa cataga cootta ga coottaat caa gag ctacca act ct ttttcc gaa gg taact gg cttcag cag gag cag a tacca a at act gt ccttct a g t g tag ccg tag gt a g ccac act ttttcc ga gag cag a gag cag a gag acaagaactotgtagcaccgcctacatacctegctetgctaatcctgttaccagtggctgctgccagtggcgataagtcgtgtcttaccgggttggactca ${\tt 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SEQUENCE ID NO. 6

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SEQUENCE ID NO. 7

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SEQUENCE ID NO. 8

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SEQUENCE ID NO. 11

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Claims

- 1. A method of identifying DNA responsible for conferring a particular phenotype in a cell which method comprises
- a) constructing a cDNA or genomic library of the DNA of said cell in a suitable vector in an orientation relative to a promoter(s) capable of initiating transcription of said cDNA or DNA to double stranded (ds) RNA upon binding of an appropriate transcription factor to said promoter(s),
- b) introducing said library into one or more of said cells comprising said transcription factor, and
- c) identifying and isolating a particular

 phenotype of said cell comprising said library and
 identifying the DNA or cDNA fragment from said library
 responsible for conferring said phenotype.
- A method according to claim 1 wherein said
 library is organised into hierarchical pools prior to step b).
 - 3. A method of assigning function to a known DNA sequence which method comprises
- 25 a) identifying a homologue(s) of said DNA sequence in a cell,
 - b) isolating the relevant DNA homologue(s) or a fragment thereof from said cell,
- c) cloning said homologue or fragment thereof
 into an appropriate vector in an orientation relative
 to a suitable promoter(s) capable of initiating
 transcription of dsRNA from said DNA homologue or
 fragment upon binding of an appropriate transcription
 factor to said promoter(s),
- 35 d) introducing said vector into said cell from

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step a) comprising said transcription factor, and

- e) identifying the phenotype of said cell compared to wild type.
- 4. A method according to any of claims 1 to 3 wherein said DNA library, homologue or fragment is cloned in a sense and an antisense direction relative to said promoter.
- 10 5. A method according to any of claims 1 to 3 wherein said DNA library, homologue or fragment is cloned between two promoters capable of producing dsRNA from said DNA library, homologue or fragment upon binding of said transcription factor to said promoters.
 - 6. A method according to any of claims 1 to 5 wherein said cell is adapted to express said transcription factor.

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- 7. A method according to any of claims 1 to 6 wherein said DNA library, homologue or fragment is constructed in a suitable vector which comprises a sequence of nucleotides encoding said transcription factor operably linked to a suitable promoter.
- 8. A method according to any of claims 1 to 6 wherein said transcription factor is encoded by a further vector independent of the vector including said DNA library, DNA homologue or fragment and which sequence encoding said transcription factor is operably linked to a suitable promoter.
- A method according to claim 7 or 8
 wherein said transcription factor comprises any of T7,

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T3 or SP6 polymerase.

- 10. A method according to claim 7 or 8 wherein said suitable promoter comprises any of let 858, SERCA, UL6, myo-2 or myo-3.
 - 11. A method according to any of claims 7 to 10, wherein said suitable vector or said further vector comprises a selectable marker.

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- 12. A method according to claim 11 wherein said selectable marker comprises a nucleotide sequence capable of inhibiting or preventing expression of a gene in said cell and which gene is responsible for conferring a known phenotype.
- 13. A method according to claim 12 wherein said nucleotide sequence comprises a sequence which is a part of or identical to said gene conferring said phenotype, and which nucleotide sequence is itself oriented relative to a suitable promoter(s) capable of initiating transcription of double stranded RNA upon binding of an appropriate transcription factor to said promoters.

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14. A method according to claim 12 wherein said nucleotide sequence is a part of or identical to said gene sequence conferring said phenotype, and which nucleotide sequence is such as to permit integration of said suitable or further vector by homologous recombination in the genome of said cell and following said integration said nucleotide sequence is capable of inhibiting expression of said gene sequence conferring said phenotype.

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15. A method according to claim 14 wherein said nucleotide sequence comprises stop codons sufficient to prevent translation of said nucleotide sequence following its integration into said genome.

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- 16. A method according to any preceding claim wherein said cell is a microorganism suitable for feeding to, transforming or infecting an organism.
- 10 17. A method according to any of claims 1 to 14 wherein said cell is contained in an organism or an embryo thereof.
- 18. A method according to any of claims 1 to 17 wherein said promoters are T7 promoters.
 - 19. A method according to any of claims 12 to 18 wherein said known gene sequence comprises a sup-35 gene or a fragment thereof which is selectable by identifying offspring growing at a temperature above 25°C following introduction of said vector in the genome of a pha-1 et123ts mutant *C. elegans* worm.
- 20. A method according to any of claims 1 to 19
 wherein said cell or organism is contacted with a
 specified compound for screening for a desired
 phenotype, such as resistance or sensitivity to said
 compound when compared to the wild type cell or
 organism.

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- 21. A method according to any preceding claim wherein said transcription factor is inducible.
- 22. A method according to claim 16 wherein said microorganism is an E. coli strain which is an RNAase

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III and preferably an RNAase negative strain.

- 23. A method according to any of claims 17 wherein said organism is selected from the class nematoda.
- 24. A method according to claim 23 wherein said organism is Caenorhabditis elegans.
- 25. A method of generating a transgenic non-human organism comprising an exogenous transcription factor and a transgene comprising a promoter operably linked to DNA fragment which is expressed upon binding of said transcription factor thereto, the method comprising
 - a) providing a first transgenic organism comprising a first construct incorporating DNA encoding an exogenous transcription factor and a second transgenic organism comprising a second construct including at least one promoter operably linked to a desired DNA sequence which is expressed upon binding of the transcription factor of said first transgenic organism thereto,
- b) crossing said first and second transgenic
 organisms and selecting offspring expressing said desired DNA sequence.
- 26. A method according to claim 25 wherein said first and second transgenic organisms are generated by transforming said first and second constructs into respective microorganisms for subsequent feeding to the respective organism.
- 27. A method according to claim 25 or 26 wherein said second construct comprises said desired DNA

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sequence in an orientation relative to said promoter so as to be capable of initiating transcription of said DNA to dsRNA upon binding of said transcription factor thereto.

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- 28. A method according to claim 27 wherein said second construct comprises two promoters flanking said desired DNA sequence which promoters can initiate transcription of said DNA sequence to dsRNA upon binding of said transcription factor to said promoters.
- DNA sequence is provided in a sense and an antisense orientation relative to said promoter so as to produce dsRNA upon binding of the transcription factor to the promoter.
- 30. A method according to any of claims 25 to 29
 wherein said second transgenic organism further
 comprises a reporter gene operably linked to a
 promoter which is capable of initiating transcription
 of said reporter upon binding of said transcription
 factor thereto.

- 31. A method according to any of claims 25 to 30 wherein said transcription factor comprises a polymerase.
- 30 32. A method according to claim 31 wherein said polymerase comprises any of T7, T3 or SP6 polymerase.
- 33. A method according to any of claims 25 to 31
 wherein said promoters comprises any of T7, T3 or SP6
 promoters.

34. A method according to claim 30 wherein said reporter gene comprises any of those sequence encoding Luciferase, Green Fluorescent protein, β galactosidase or $\beta\text{-lactamase}.$

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- 35. A method according to any of claims 25 to 30 wherein said organism is of the species nematoda.
- 36. A method according to claim 35 wherein said nematoda species is *C. elegans*.
 - 37. A transgenic non-human multicellular organism obtainable according to the methods of any one of claims 25 to 34.

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- 38. A method of validating clones identified in yeast two hybrid vector experiments which method comprises
- a) providing a construct including the DNA

 20 encoding the protein identified in the two hybrid
 vector experiment, which construct is such that said
 DNA is orientated relative to a promoter(s) that is
 capable of initiating transcription of said DNA to
 double stranded RNA upon binding of an appropriate
 transcription factor to said promoter(s),
 - b) transforming a cell comprising said transcription factor with said construct, and
 - c) identifying a phenotypic change in said cell or organism when compared to a wild type.

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39. A method according to claim 38 wherein said DNA sequence is provided between two promoters capable of initiating transcription of the DNA sequence to dsRNA upon binding of the transcription factor to

said promoters.

- 40.A method according to claim 38 wherein said DNA is provided in a sense and an antisense orientation relative to said promoter such that binding of the transcription factor to said promoter initiates transcription of dsRNA from said DNA.
- 41. A method according to any of claims 38 to 40
 wherein said transcription factor is inducible in said cell.
 - 42. A method according to any of claims 38 to 41 wherein said promoter is a phage polymerase promoter and said transcription factor is a RNA polymerase.
 - 43. A method according to claim 42 wherein said polymerase is any of T7 RNA polymerase, T3 RNA polymerase or SP6 RNA polymerase.

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- 44. A method according to claim 43 wherein said promoters comprise any of T7, T3 or SP6 promoter
- 45. A method according to any of claims 38 to 44
 25 wherein said construct is such that it may be used in yeast two hybrid experiments.
 - 46. A method according to any of claims 38 to 45 wherein said cell is an $E.\ coli$ cell.

- 47. A method according to any of claims 38 to 45 wherein said cell is part of an organism or an embryo thereof.
- 35 48. A method according to claim 47 wherein said

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organism is of the species nematoda and preferably C. elegans.

49. Plasmid pGN1 as illustrated in Figure 1.

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- 50. Plasmid pGN100 as illustrated in Figure 2.
- 51. The yeast two hybrid vector plasmid illustrated in any of Figures 4, 15 or 16(Seq ID Nos 8 and 9).
 - 52. A plasmid as illustrated in Figure 7.
 - 53. A plasmid as illustrated in Figure 8.

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- 54. A method of alleviating infestation of plant pests, which method comprises
- a) identifying a DNA sequence from said pest which is critical for its survival, growth, proliferation,
- b) cloning said sequence from step a) or a fragment thereof in a suitable vector in an orientation relative to promoter(s) such that said promoter(s) is capable of initiating transcription of said DNA sequence to RNA or dsRNA upon binding of an appropriate transcription factor to said promoter(s), and
 - c) introducing said vector into the plant.
- DNA sequence is provided between two promoters such that binding of the transcription factor to the promoters results in transcription of the DNA to dsRNA.

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- 56. A method according to claim 54 wherein said DNA sequence is provided in a sense and an antisense orientation relative to said promoter such that binding of the transcription factor to the promoter results in transcription of the DNA to dsRNA.
- 57. A method according to claim 54 wherein said pest is a nematode worm.
- 10 58. A method according to claim 57 wherein said nematode comprises any of Tylenchulus ssp. Radopholus ssp., Rhadinaphelenchus ssp., Heterodera ssp., Rotylenchulus ssp., Pratylenchus ssp., Belonolaimus ssp., Canjanus ssp., Meloidogyne ssp., Globodera ssp., Nacobbus ssp., Ditylenchus ssp., Aphelenchoides ssp., Hirschmenniella ssp., Anguina ssp., Hoplolaimus ssp., Heliotylenchus ssp., Criconemellassp., Xiphinemassp., Longidorus ssp., Trichodorus ssp., Paratrichodorus ssp., Aphelenchs ssp.

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- 59. A method according to claim 55 wherein said DNA sequence or fragment thereof is cloned between two tissue, preferably root specific promoters.
- 25 60. An expression vector for use in a method according to any proceeding claim comprising a promoter or promoters oriented relative to a DNA sequence such that they are capable of initiating transcription of said DNA sequence to double stranded RNA upon binding of an appropriate transcription factor to said promoter or promoters.
- 61. An expression vector according to claim 60 comprising two identical promoters flanking the DNA sequence.

62. An expression vector according to claim 60 comprising said DNA sequence in a sense and an antisense orientation relative to said promoter.

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- 63. An expression vector according to any of claims 60 to 62 which further comprises a nucleotide sequence encoding a selectable marker.
- wherein said nucleotide sequence encoding said selectable marker is orientated relative to the promoter(s) such that transcription of the nucleotide sequence to double stranded RNA occurs upon binding of an appropriate transcription factor to said promoter(s).
- wherein said nucleotide sequence encoding the selectable marker is provided between the identical promoters capable of initiating transcription of the nucleotide sequence to dsRNA upon binding of the transcription factor to the promoters.
- wherein said nucleotide sequence encoding the selectable marker is provided in a sense and an antisense orientation relative to the promoter such that transcription of the nucleotide sequence to dsRNA upon binding of the transcription factor to said promoter occurs.
 - 67. An expression vector according to claim 63 or 64 wherein said selectable marker comprises a nucleotide sequence encoding sup-35, for introduction into *C. elegans* having a pha-1 mutation.

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- 68. An expression vector for expressing an appropriate transcription factor for use in a method according to any of claims 1 to 48 and 54 to 59 which vector comprises a sequence of nucleotides encoding said transcription factor operably linked to suitable expression control sequences.
- 69. An expression vector according to claim 68 wherein said expression control sequences include promoters which are inducible, constitutive, general or tissue specific promoters, or combinations thereof.
- 70. An expression vector according to any of claims 68 to 69 wherein said transcription factor comprises a phage polymerase, and preferably T7 RNA polymerase.
- 71. An organism or cell transformed or transfected with a plasmid according to any of claims 49 to 53 or an expression vector according to any of claims 60 to 70.
 - 72. An organism according to claim 71, which is of the species nematoda and preferentially C. elegans.

73. A method of introducing dsRNA or DNA capable of producing dsRNA into an organism which method comprises feeding said organism with a suitable microorganism comprising an expression vector according to any of claims 60 to 67 or feeding said

- organism directly with an expression vector according to any of claims 60 to 67.
- 74. A method according to claim 73 wherein said35 microorganism or said organism is adapted to express

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said transcription factor.

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- 75. A method according to claim 74 wherein either said microorganism or said organism comprises an expressing vector according to any of claims 66 to 69.
- 76. A method according to any of claims 73 to 75 wherein said organism is *C. elegans* and said microorganism is *E. coli*.
 - 77. A method according to claim 76 wherein said $\it E.~coli$ strain is an RNAaseIII negative strain.
- 78. A method according to any of claims 73 to 75 wherein said organism is a *C. elegans* nuc-1 mutant.
- 79. A selection system for identifying transformation of a cell or organism with a vector according to claims 60 to 63 which system comprises a vector according to claims 60 to 63 and said selectable marker comprises a nucleotide sequence capable of inhibiting or preventing expression of a gene in said cell or organism which gene is responsible for conferring a known phenotype.
- wherein said nucleotide sequence comprises a sequence which is a part of or identical to said gene

 conferring said known phenotype, and which nucleotide sequence is itself located between two identical promoters capable of initiating transcription of double stranded RNA upon binding of an appropriate transcription factor to said promoters.

wherein said nucleotide sequence is a part of or identical to said gene sequence which confers a known phenotype on said cell or organism, and which nucleotide sequence permits integration of said vector by homologous recombination in the chromosome of said cell or organism and following said integration said sequence inhibits expression of said gene sequence conferring said known phenotype.

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82. A selection system according to claim 81 wherein said nucleotide sequence comprises stop codons sufficient to prevent translation of said nucleotide sequence following integration into said genome.

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- 83. A selection system according to claim 79 wherein said known gene sequence comprises a sup-35 gene or a fragment thereof which is selectable by identifying offspring growing at a temperature above 25°C following introduction of said vector in a pha-1 et123ts mutant *C. elegans* worm.
- 84. A method according to claim 74 wherein said transcription factor is T7 RNA polymerase.

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- 85. A method of assigning function to a DNA sequence of a multicellular organism which method comprises:
 - a) providing:
- (i) a first construct comprising said DNA sequence cloned in a sense direction under the regulation of a suitable promoter
 - (ii) a second construct comprising said DNA sequence cloned in an anti-sense direction under the regulation of the same promoter as in step (i),

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in a multicellular organism which is adapted to initiate transcription of said DNA fragment from said promoter,

- b) identifying the phenotype of said
 multicellular organism compared to wild type.
 - 86. A method of assigning function to a DNA sequence of a multicellular organism which method comprises:
- 10 a) providing
 - i) a construct comprising said DNA fragment cloned between two promoters capable of promoting transcription in said multicellular organism,

in a multicellular organism capable of initiating transcription from said promoter;

- b) identifying the phenotype of said multicellular organism compared to wild type.
- 87. A method for expressing a gene, cDNA or DNA fragment in C. elegans which method comprises :
 - i) providing a transgenic C. elegans expressing an exogenous transcription factor,
 - ii) cloning said gene or cDNA or DNA fragment into an appropriate vector operably linked to a promoter capable of initiating transcription of said gene, cDNA or DNA fragment upon binding of said exogenous transcription factor thereto,
 - iii) introducing said vector into said transgenic C. elegans from step i).

88. A method according to claim 87 wherein said exogenous transcription factor is a phage polymerase.

89. A method according to claim 88 wherein

said phage polymerase is T3, SP6 or T7 RNA polymerase.

90. A method according to any of claims 87 to 89 wherein said promoter is any of T3, T7 or SP6 promoter.

F1G. 1.

pGNI

gagtgcaccatatgcggtgtgaaataccgcacagatgcgtaaggagaaaataccgcatcaggcgaaattgtaaacgttaatattt tgttaaaattcgcgttaaatatttgttaaatcagctcattttttaaccaataggccgaaatcggcaaaatcccttataaatcaaaagaatagaccgagatagggttgagtgttccagtttggaacaagagtccactattaaagaacgtggactccaacgtcaaagggcgaa aaaccgtctatcagggcgatggcccactacgtgaaccatcacccaaatcaagttttttgcggtcgaggtgccgtaaagctctaaat cgaaaggagcgggcgctagggcgctggcaagtgtagcggtcacgctgcgcgtaaccaccacacccgccgcgcttaatgcgc egetacagggegegtecanegecaneaggetgegeaactgngggaagggegateggngegggeetenegetanaegeca gctggcgaaagggggatgtgctgcaaggcgattaagttgggtaacgccagggttttcccagtcacgacgttgtaaaacgacgg ccagtgaattgtaatacgactcactatagggcgaattcgagctcggtacccgggggatcctctagagtcgaaagcttctcgccctat agtgagtcgtattacagcttgagtattctatagtgtcacctaaatagcttggcgtaatcatggtcatagctgtttcctgtgtgaaattgtt aattgcgttgcgctcactgcccgctttccagtcgggaaacctgtcgtgccagctgcattaatgaatcggccaacgcgcggggag aggeggfttgegtattgggegetetteegetteetegeteactgactegetgegeteggtegtteggetgeggeggeggtateag ctcactcaaaggcggtaatacggttatccacagaatcaggggataacgcaggaaagaacatgtgagcaaaaaggccagcaaaa ggccaggaaccgtaaaaaaggccgcgttgctggcgtttttcgataggctccgccccctgacgagcatcacaaaaatcgacgct caagtcagaggtggcgaaacccgacaggactataaagataccaggcgtttccccctgggaagctccctcgtgcgctctctgttc cgaccctgccgcttaccggatacctgtccgcctttctcccttcgggaagcgtggggctttctcatagctcacgctgtaggtatctca tegtettgagtecaaeeeggtaagaeaegaettategeeaetggeageageeaetggtaaeaggattageagagegaggtatgt aggcggtgctacagagttcttgaagtggtggcctaactacggctacactagaaggacagtatttggtatctgcgctctgctgaagc cagitacciteggaaaaagagitggtagcictigateeggeaaacaaaccacegetggtageggitggittitigtitgcaagcage agattacgcgcagaaaaaaaggatctcaagaagatcctttgatcttttctacggggtctgacgctcagtggaacgaaaactcacgt atgagtaaacttggtctgacagttaccaatgcttaatcagtgaggcacctatctcagcgatctgtctatttcgttcatccatagttgcct gacteceegtegtgtagataactaegataegggagggettaeeatetggeeecagtgetgeaatgataeegegagaeeeaeget

FIG. 1. (CONTINUED)

FIG. 2.

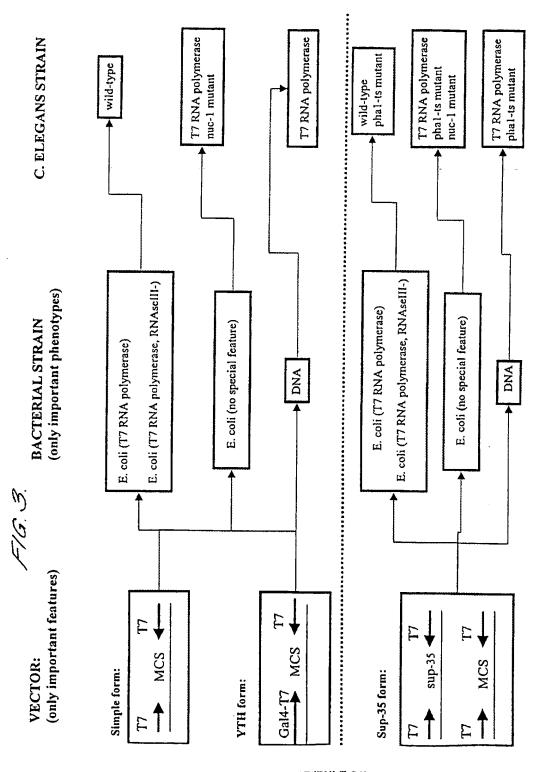
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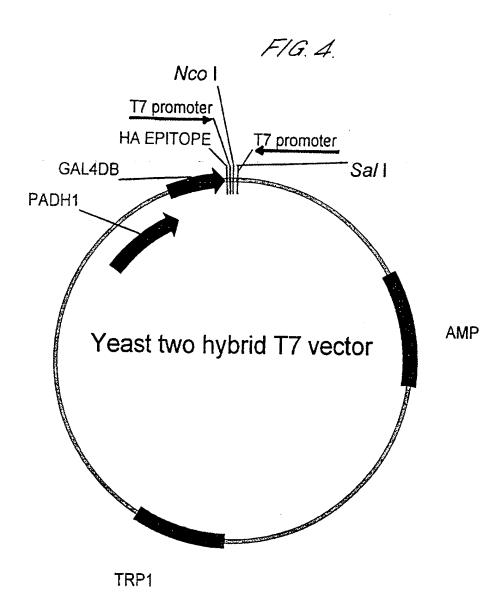
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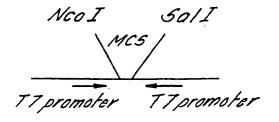
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FIG. 2. (CONTINUED 2)



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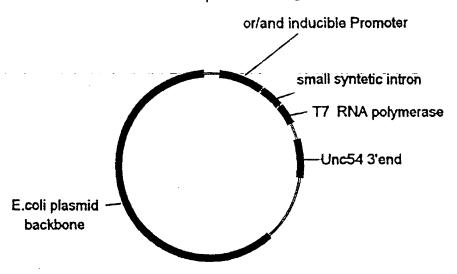


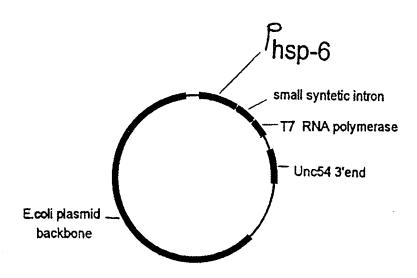
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FIG. 5. 7/21

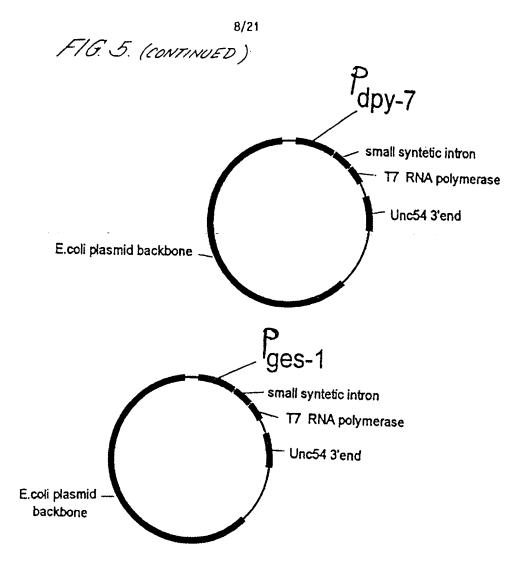
General description of the C.elegans T7 RNA polymerase expression vector with 4 examples

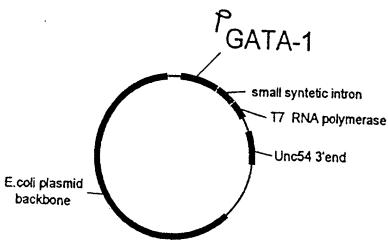
Specific or/and general or/and constitutive





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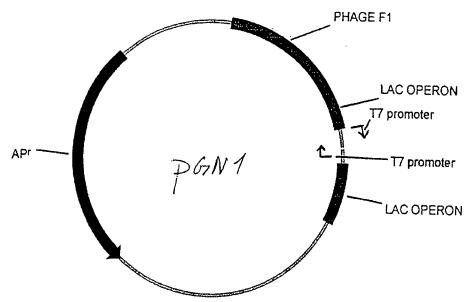


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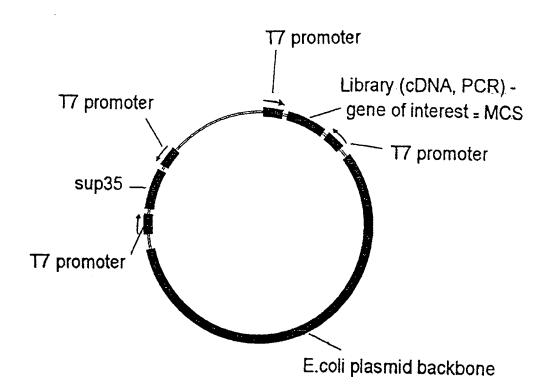
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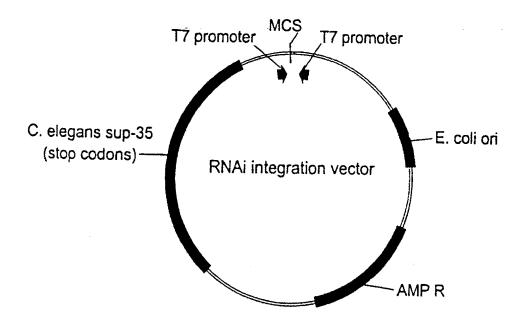


F1G. 7.



enhanced vector for RNAi, producing sup 35 ds RNA and ds RNA of the library, gene of interest or PCR product.

F16. 8.



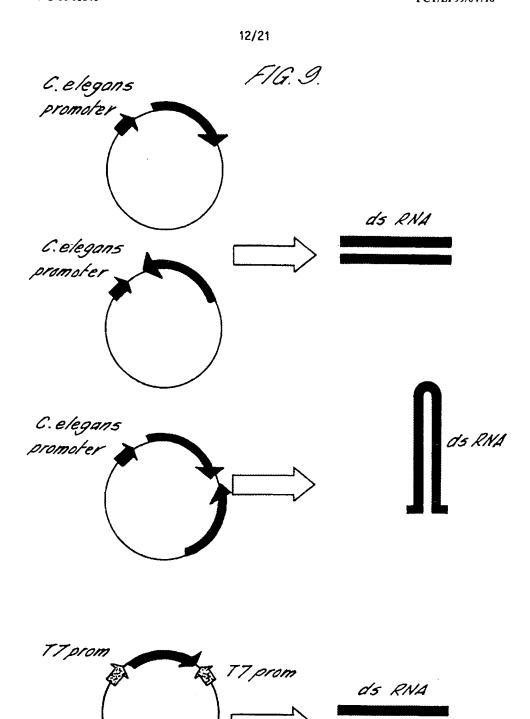
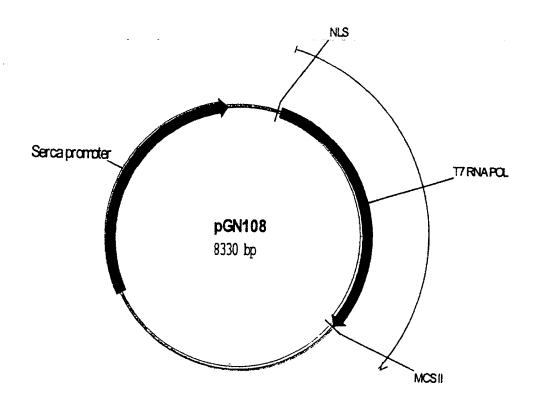
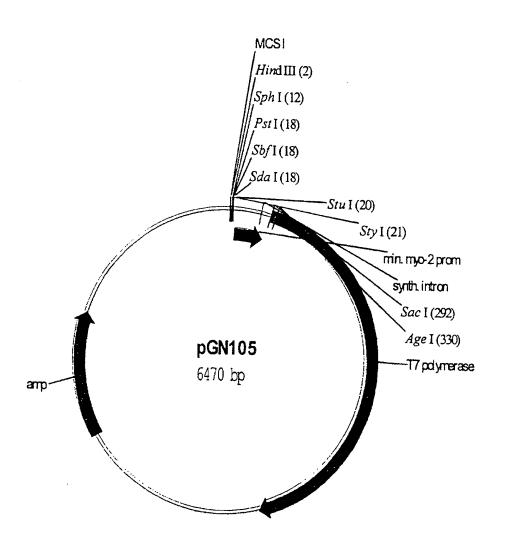


FIG. 10.

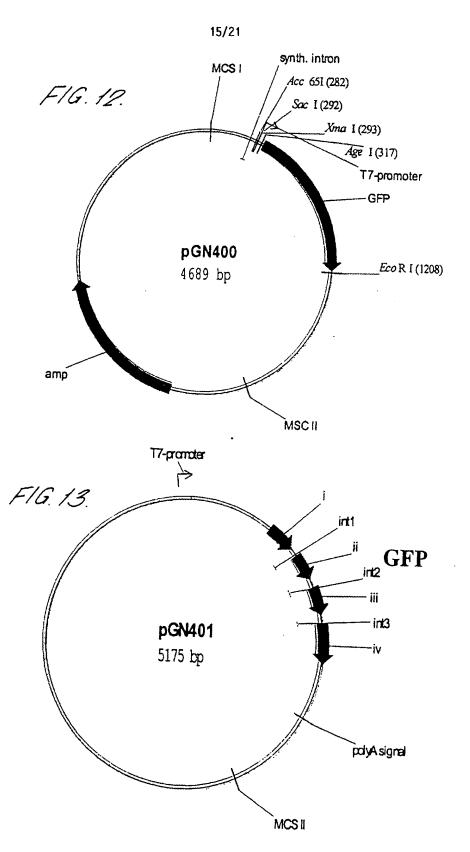






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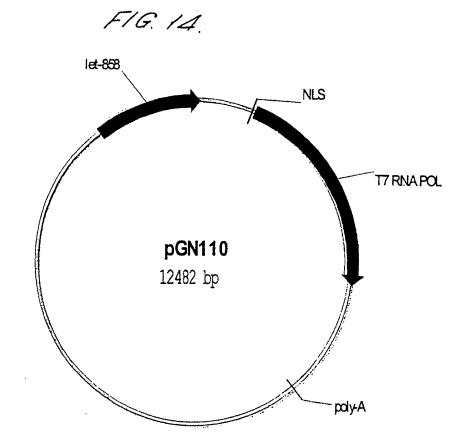
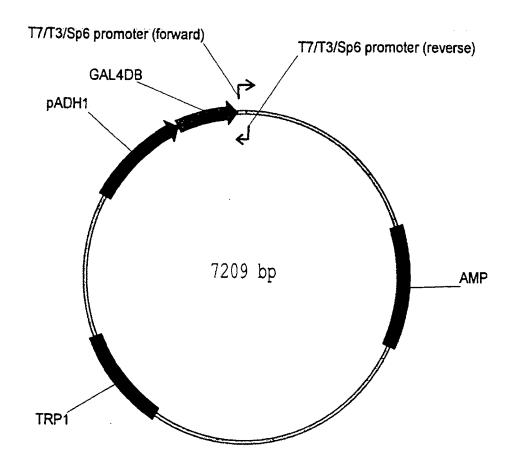


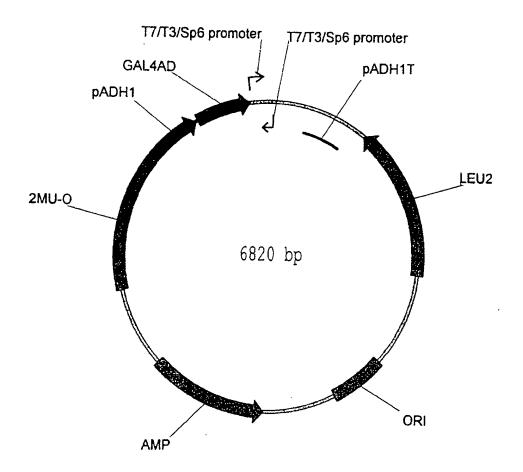
FIG. 15.

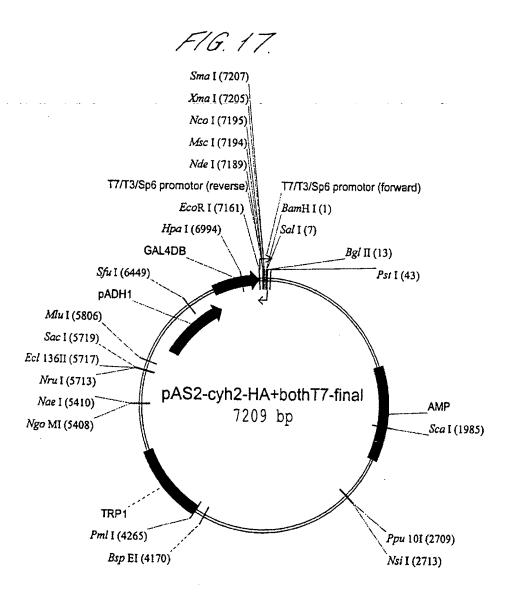
pAS2* with Forward and Reverse T7/T3/Sp6

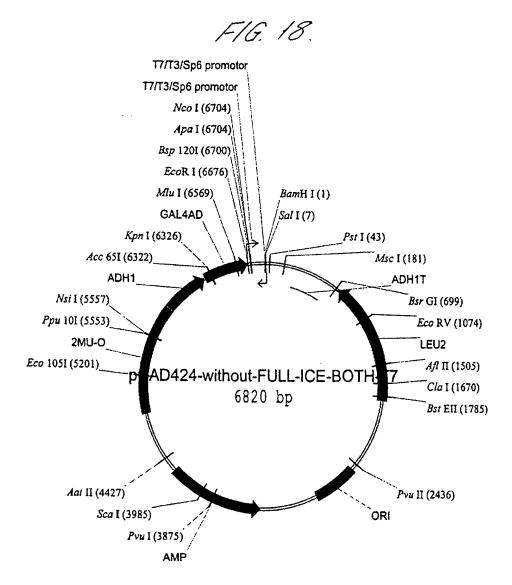


F1G. 16.

pGAD424 with Forward and Reverse T7/T3/Sp6

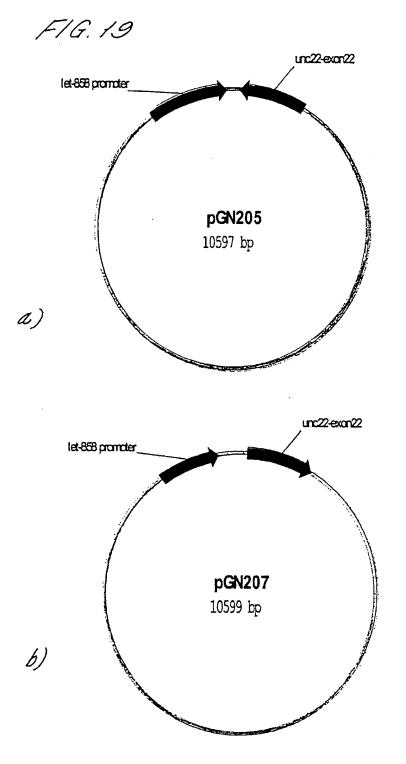






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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 7; C12Q 1/68, C12N 15/63	A3	(11) International Publication Number: WO 00/01846
(21) International Application Number: PCT/EP (22) International Filing Date: 2 July 1999 ((30) Priority Data: 9814536.0 3 July 1998 (03.07.98) 9827152.1 9 December 1998 (09.12.98) (71) Applicant (for all designated States except US): DEVC [BE/BE]; Technologiepark 9, B-9052 Wondelgem (72) Inventors; and (75) Inventors/Applicants (for US only): PLAETINC [BE/BE]; Devgen N.V., Technologiepark 9, Wondelgem (BE), PLATTEEUW, Christ [BE/BE] N.V., Technologiepark 9, B-9052 Wondelgem MORTIER, Katherine [BE/BE]; Devgen N.V., Technologiepark 9, Wondelgem (BE). BOGAERT [BE/BE]; Devgen N.V., Technologiepark 9, Wondelgem (BE). (74) Agent: BOULT WADE TENNANT; 27 Furniva London EC4A IPQ (GB).	O2.07.9 COLOT.9 COL	BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published With international search report. (88) Date of publication of the international search report: 15 June 2000 (15.06.00)

(54) Title: CHARACTERISATION OF GENE FUNCTION USING DOUBLE STRANDED RNA INHIBITION

(57) Abstract

There is provided a method of identifying DNA responsible for conferring a particular phenotype in a cell which method comprises a) constructing a cDNA or genomic library of the DNA of said cell in a suitable vector in an orientation relative to a promoter(s) capable of initiating transcription of said cDNA or DNA to double stranded (ds) RNA upon binding of an appropriate transcription factor to said promoter(s), b) introducing said library into one or more of said cells comprising said transcription factor, and c) identifying and isolating a particular phenotype of said cell comprising said library and identifying the DNA or cDNA fragment from said library responsible for conferring said phenotype. Using this technique it is also possible to assign function to a known DNA sequence by a) identifying a homologue(s) of said DNA sequence in a cell, b) isolating the relevant DNA homologue(s) or a fragment thereof from said cell, c) cloning said homologue or fragment thereof into an appropriate vector in an orientation relative to a suitable promoter(s) capable of initiating transcription of dsRNA from said DNA homologue or fragment upon binding of an appropriate transcription factor to said promoter(s) and d) introducing said vector into said cell from step a) comprising said transcription factor.

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inte onal Application No

		l P	CT/EP 99/04718
A. CLASS	IFICATION OF SUBJECT MATTER C12Q1/68 C12N15/63	·····	
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IPC 7	C12Q C12N	and symbols;	
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Electronic d	lata base consulted during the international search (name of data	base and, where practical sea	rch (erms used)
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C. DOCUM	ENTS CONSIDERED TO BE RELEVANT		· · · · · · · · · · · · · · · · · · ·
Category *	Citation of document, with indication, where appropriate, of the r	relevant passages	Relevant to claim No.
Χ	WO 90 14090 A (HEM RES INC) 29 November 1990 (1990-11-29)		60,62,63
	page 5, line 2 -page 6, line 18		

X	WO 91 15111 A (HARVARD COLLEGE) 17 October 1991 (1991-10-17)		25,37
Υ	page 2, line 20 -page 13, line 1	l; claims	87-90
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<u> </u>	er documents are bated in the continuation of box C.	X Patent lamily memb	pers ara listed in annex.
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29	March 2000	714. 04.	00
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Inte onal Application No PCT/EP 99/04718

C.(Continu	ation) DOCUMENTS CONSIDERED TO BE RELEVANT	PCT/EP 99/04718
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Х	VENNEMA H ET AL: "Enhancement of the vaccinia virus/ phage T7 RNA polymerase expression system using encephalomyocarditis virus 5'-untranslated region sequences." GENE, (1991 DEC 15) 108 (2) 201-9., XP000872841 page 1	70,71
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Y	page 11, line 9 - line 34; claims 26,27 FIRE A ET AL: "Potent and specific genetic interference by double-stranded RNA in Caenorhabditis elegans" NATURE,GB,MACMILLAN JOURNALS LTD. LONDON, vol. 391, 19 February 1998 (1998-02-19), pages 806-811, XP002095876 ISSN: 0028-0836 cited in the application the whole document	1-24, 38-48, 85,86
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A	DATABASE MEDLINE 'Online! US NATIONAL LIBRARY OF MEDICINE (NLM), BETHESDA, MD, US JAMES P ET AL: "Genomic libraries and a host strain designed for highly efficient two-hybrid selection in yeast." retrieved from STN Database accession no. 97132579 XP002130070	51
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4

International application No. PCT/EP 99/04718

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)
This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:
Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:
2. X Claims Nos.: 49,50,68,69,79-83 because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
see FURTHER INFORMATION sheet PCT/ISA/210
Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).
Box II Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)
This International Searching Authority found multiple inventions in this international application, as follows:
see additional sheet
As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims
As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. No required additional search fees were timely paid by the applicant. Consequently, this international Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:
Remark on Protest The additional search fees were accompanied by the applicant's protest. X No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box 1.2

Claims Nos.: 49,50,68,69,79-83

- 1) Claims 49 and 50 relate to sequences which have been filed as figures, i.e. not in a searchable format.
- 2) Claims 68 and 69 do not relate to any technical feature which would allow a meaningful search to be carried out because the expression "appropriate transcription factor for use in a method according to claims..." is too vague and devoid of a clear technical meaning.
- 3) Claim 79 relates to a "selection system" which is solely characterised by comprising the vector of claims 60-63; no further features of the claimed system are apparent: its subject-matter is thus the same as that of the said claims 60-63. No additional features are present which could be the basis of the search: hence, a search as to the subject-matter of claim 79 and 80-84 dependent thereon is not feasible.

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

1. Claims: 1-24,27-29, 30-37 (part),48,51-67,70-78,85,86

Methods for identifying a DNA responsible for a particular phenotype, for assigning function to a known DNA sequence, for generating a transgenic animal, for alleviating plant pests infestation, for introducing dsRNA into an organism, plasmids, expression vectors, transformed cells

2. Claims: 25,26,30-37 (part),87-90

Method for obtaining a transgenic animal and method for expressing a gene in a transgenic C. elegans

information on patent family members

Inte 'onal Application No PCT/EP 99/04718

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	atent document d in search repo		Publication date		Patent family member(s)	Publication date
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